



EU declaration of conformity

| | | |
|---|---|---------------|
| Manufacturer according to Regulation 2017/745 | Schülke & Mayr GmbH Robert-Koch-Str. 2 22851 Norderstedt Germany | |
| Registration Number acc. to Art. 31 2017/745 | DE-MF-000005701 | |
| Product name | thermosept® X-tra | |
| basic UDI-DI | 4032651-BSC00000004-CT | |
| Code acc. to Art. 26 2017/745 | V0799 | |
| Intended Purpose | cleaning agent for automated reprocessing of medical devices | |
| Risk Class according to Regulation 2017/745 | I | |
| | annex | VIII |
| | rule | 1 |
| Standards applied | EN ISO 13485 additional standards see technical documentation Schülke & Mayr GmbH | |
| Conformity Assessment Procedure according to Regulation 2017/745 | annex | IV / V |
| Certificate | EN ISO 13485 | 004567 MP2016 |
| Version | 2-0 | |

Schülke & Mayr GmbH herewith declares that the device covered by this declaration is in conformity with the Regulation 2017/745 concerning medical devices.

Schülke & Mayr GmbH declares that Schülke & Mayr GmbH bears the sole responsibility for issuing this Declaration

Norderstedt **09. Nov. 2021**
ppa.


Dr. Uwe Berlekamp
Schülke & Mayr GmbH
Director Business Line Healthcare

ppa.

Jörn Ahlsdorff
Schülke & Mayr GmbH
Director Industrial Operations
International Industrial Operations